Claims

pharmaceutical preparation of oxaliplatinum in liquid form, wherein said flexible hag is constructed from plastic material, excluding, for the said material in direct contact with the said pharmaceutical preparation, PVC-based material.

- 2) Flexible impervious bag for medical use containing a pharmaceutical preparation of oxaliplatinum according to Claim 1, wherein the envelope of said bag has a multi-layer structure.
- 3) Flexible impervious bag for medical use containing a pharmaceutical preparation of oxaliplatinum according to one of the foregoing Claims, wherein the liquid solution of oxaliplatinum is in contact with an internal layer of the said envelope consisting of a polypropylene material.
- 4) Flexible impervious bag for medical use containing a pharmaceutical preparation of oxaliplatinum according to one of the foregoing Claims, wherein the concentration of oxaliplatinum in the pharmaceutical preparation is between 1 and 8 mg/ml.
- 5) Flexible impervious bag for medical use containing a pharmaceutical preparation of oxaliplatinum according to Claim 4, wherein said concentration is between 1 and 5 mg/ml.
- 6) Flexible impervious bag for medical use containing a pharmaceutical preparation of oxaliplatinum according to one of the foregoing Claims, wherein said the file bag consists of two welded sheets of multi-layer sheet material comprising one film of polyamide of 11-amino-undecanoic acid bonded by at least one of its surfaces to a film of polypropylene by means of a film of polyolefine, the polypropylene films forming the internal wall of the watertight flexible bag.



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- pharmaceutical preparation of oxaliplatinum according to Claim 7, wherein the multi-compartments are defined in such a way as to allow the dosing of a ready-for-use preparation.
- 9) Flexible impervious bag for medical use containing a pharmaceutical preparation of oxaliplatinum according to one of the foregoing Claims, wherein said solution has a pH of 4.5 to 6.0, a concentration of oxaliplatinum in the preparation of at least 95 % of the initial concentration, as well as a clear, colourless appearance free from precipitate after storage for a pharmaceutically acceptable period.

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AMENDED PAGE